EXHIBIT 11

to

PAUL D. BRACHMAN DECLARATION IN SUPPORT OF DEFENDANT'S TRIAL BRIEF

ĺ	Case 3:21-cv-03496-AMO	Document 405-6	Filed 01/08/25	Page 2 of 9				
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7 8	Attorneys for Plaintiff and Counter-Defendant, SURGICAL INSTRUMENT SERVICE COMPANY, INC.							
9	UNITED STATES DISTRICT COURT							
10	NORTHERN DISTRICT OF CALIFORNIA							
11	SAN FRANCISCO DIVISION							
12	CLID CLC AL DICTRIA (T)	TT CEDY HOE	G N 221 0	2407 4240				
13	SURGICAL INSTRUMEN COMPANY, INC.	NT SERVICE	Case No. 3:21-cv-03 Honorable Araceli N					
14	Plaintiff/Counter-Defendant,		PLAINTIFF SIS'S RESPONSES TO					
15	V.		DEFENDANT INTUITIVE SURGICAL, INC.'S SECOND SET OF REQUESTS FOR ADMISSION AND THIRD SET OF					
16	INTUITIVE SURGICAL, INC.,		INTERROGATORIES					
17	Defendant/Counterclaimant.							
18	PROPOUNDING PARTY:	Defendant Intuitive	Surgical, Inc.					
19	RESPONDING PARTY:							
20 21	SET NUMBER:							
22	SET NOMBER.	Two of Requests for Admission						
23	Three of Interrogatories							
24	Pursuant to Rules 26 and 36 of the Federal Rules of Civil Procedure Plaintiff Surgical							
25	Instrument Service Co., Inc. ("Plaintiff" or "SIS"), by and through the undersigned counsel, hereby							
26	responds to Defendant Intuitive Surgical, Inc.'s ("Defendant" or "Intuitive") Second Set of							
27	Requests for Admission and Third Set of Interrogatories as follows:							
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	SIS's RESPONSES TO 2ND RFA AND 3RD INTERROGATORIES 3:21-cv-03496-AMO							

GENERAL OBJECTIONS

Plaintiff makes the following general responses and objections ("General Objections") to each definition, instruction, and request propounded in Defendant's second set of requests for admission. These General Objections are hereby incorporated into each specific response. The assertion of the same, similar, or additional objections or partial responses to individual requests does not waive any of Plaintiff's General Objections.

Plaintiff reserves its right to and will supplement its responses to Defendant's second set of requests for admission in accordance with Fed. R. Civ. P. 26(e) and the objections as set forth herein are made without prejudice to Plaintiff's right to assert any additional or supplemental objections pursuant to Fed. R. Civ. P. 26(e).

- 1. Plaintiff objects to Defendant's definition of "EndoWrist Instruments" as vague and ambiguous, and further on the basis that determining the scope of "Intuitive's patented 'EndoWrist' technology" necessarily requires legal opinion and testimony.
- 2. Plaintiff objects to the definitions and instructions accompanying Defendant's Requests to the extent they seek to impose duties or obligations upon Plaintiff greater than required by the Federal Rules of Civil Procedure and Local Rules for the Northern District of California.
- 3. Plaintiff objects to each request to the extent that it seeks information that is protected from disclosure by the attorney-client privilege, the attorney work product doctrine or any other applicable privilege, doctrine, or discovery immunity. Plaintiff's answers do not waive any attorney client privilege and Defendant may not use Plaintiff's answers to support an argument or a further line of questioning regarding the legal conclusion a request calls for.
- 4. Plaintiff objects to Defendant's requests to the extent that they are vague, ambiguous, or seek to impose obligations on Plaintiff that are broader than, or inconsistent with,

the Federal Rules of Civil Procedure or the local rules of this Court. Unless indicated otherwise, Plaintiff shall give the terms of these requests their ordinary and plain meanings. If Defendant subsequently assert an interpretation of any request that differs from Plaintiff's understanding, Plaintiff reserves the right to amend or supplement its objections and/or responses.

5. Plaintiff objects to Defendant's definition of "Customer" as vague and ambiguous. Plaintiff objects to "Customer" including any person who was "marketed" any "SIS services," as such definition is vague, ambiguous, and potentially covers any person who has seen marketing materials of Plaintiff, without regard to whether Plaintiff has engaged with or even knows of such person. Plaintiff further objects to Defendant's definition of "Customer" as including any person who has "inquired about" any "SIS services," as such definition is vague and ambiguous, for example, by lacking any limitation as to whether such person has made an inquiry to Plaintiff, whether Plaintiff has any knowledge of such inquiry, or whether such inquiry is made for the purpose of potentially purchasing SIS services. Plaintiff further objects to "Customers" as being vague and ambiguous in view of Plaintiff and Defendant both being in a market in which the party making a purchasing decision may range from individual surgeons or surgical centers, to hospitals, to entire hospital systems or purchasing organizations.

SPECIFIC OBJECTIONS AND RESPONSES

REQUEST FOR ADMISSION NO. 16: Admit that SIS did not, after November 2022, whether alone or with a third party, seek FDA clearance for any service, procedure, or technology for resetting or reprogramming the use counter on any EndoWrist Instrument.

Response:

Plaintiff objects to this Request because it implies that FDA clearance is required for any service, procedure, or technology for resetting or reprogramming the use counter on any EndoWrist

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Instrument. That is not the case. As the Court stated in the Order Re: Cross Motions for Summary Judgment (Doc. 204 p. 10):

"[T]his query is problematic because '[t]he FDCA [(Food, Drug and Cosmetic Act)] leaves no doubt that it is the Federal Government rather than private litigants who are authorized to file suit for noncompliance with the medical device provisions.' Buckman Co. v. Plaintiffs' Legal Comm., 531 U.S. 341, 349 n.4 (2001). Indeed, a private action brought under the Lanham Act may not be pursued when it requires litigating an alleged underlying FDCA violation where the FDA has not itself concluded that a violation exists. *PhotoMedex*, *Inc. v. Irwin*, 601 F.3d 919, 924 (9th Cir. 2010) (affirming grant of summary judgment for defendant on Lanham Act false advertising claim where FDA had not taken a position on defendant's laser's need for Section 510(k) clearance, and claim was based on defendant allegedly misrepresenting that its laser had received FDA clearance). * * * [C]ourts have consistently precluded private actions which require establishing a violation of the FDCA. See, e.g., Amarin Pharma, Inc. v. International Trade Commission, 923 F.3d 959, 968 (Fed. Cir. 2019) (citing *PhotoMedex*, 601 F.3d at 924, 928) (finding that Lanham Act claim could not stand where it was "based on proving violations of the FDCA and where the FDA has not taken the position that the articles at issue do, indeed, violate the FDCA.")."

The Court ruled that "SIS has not clearly engaged in unlawful conduct and accordingly may still seek to prove that Intuitive's anticompetitive conduct caused its antitrust injury at trial." Doc. 204 p. 16.

Subject to the foregoing objection, Admitted.

No Response Required

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SIS's RESPONSES TO 2ND RFA AND 3RD INTERROGATORIES 3:21-cv-03496-AMO

1	Case 3:21-cv-03496-AMO	Document 405-6	Filed 01/08/25	Page 7 of 9		
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